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FILE LAST UPDATED: 27 Apr 2011 (20110427/ED)
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=> s (transformation (w) cassette#)/bi,ab 439987

TRANSFORMATION/BI 293839

TRANSFORMATION/AB 32137 CASSETTE#/BI

24994 CASSETTE#/AB

L1 20 (TRANSFORMATION (W) CASSETTE#)/BI,AB

=> s I1 not 2011/py 600819 2011/PY

L2 18 L1 NOT 2011/PY

=> s l2 not 2010/py 1933915 2010/PY L3 12 L2 NOT 2010/PY

=> s l3 not 2009/py 1912317 2009/PY

L4 9 L3 NOT 2009/PY

=> s |4 not 2008/py 1810285 2008/PY

L5 6 L4 NOT 2008/PY

=> s I5 not 2007/py 1729373 2007/PY

L6 6 L5 NOT 2007/PY

=> s l6 not 2006/py 1591787 2006/PY

L7 6 L6 NOT 2006/PY

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L2 18 S L1 NOT 2011/PY

L3 12 S L2 NOT 2010/PY

L4 9 S L3 NOT 2009/PY

L5 6 S L4 NOT 2008/PY L6 6 S L5 NOT 2007/PY

L7 6 S L6 NOT 2006/PY

=> d l7 1-6 bib ab

L7 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2011 ACS on STN

AN 2004:593127 CAPLUS << LOGINID::20110428>>

DN 141:289470

TI VIGS vectors for gene silencing: Many targets, many tools

AU Robertson, Dominique

CS Departments of Botany and Genetics, North Carolina State University, Raleigh, NC, 27695-7612, USA

SO Annual Review of Plant Biology (2004), 55, 495-519 CODEN: ARPBDW

PB Annual Reviews Inc.

DT Journal; General Review

LA English

AB A review. The discovery that plants recognize and degrade invading viral RNA caused a paradigm shift in our understanding of viral/host interactions. Combined with the discovery that plants cosuppress their own genes if they are transformed with homologous transgenes, new models for both plant intercellular communication and viral defense have emerged. Plant biologists adapted homol.-based defense mechanisms triggered by incoming viruses to target individual genes for silencing in a process called virus-induced gene silencing (VIGS). Both VIGS- and dsRNA-contg.

transformation ***cassettes*** are increasingly being used for reverse genetics as part of an integrated approach to detg. gene function. Virus-derived vectors silence

gene expression without transformation and selection. However, because viruses also alter gene expression in their host, the process of VIGS must be understood. This review examines how DNA and RNA viruses have been modified to silence plant gene expression. Advantages and disadvantages of VIGS in detg. gene function has been discussed and guidelines for the safe use of viral vectors.

OSC.G 92 THERE ARE 92 CAPLUS RECORDS THAT CITE THIS RECORD (92 CITINGS)

RE CNT 161 THERE ARE 161 CITED REFERENCES
AVAILABLE FOR THIS RECORD ALL CITATIONS
AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2011 ACS on STN AN 2001:122995 CAPLUS < LOGINI D::20110428>> DN 136:113372

TI Chloroplast transformation in Euglena gracilis: splicing of a group III twintron transcribed from a transgenic psbK operon

AU Doetsch, Natalie A.; Favreau, Mitchell R.; Kuscuoglu, Nesrin; Thompson, Michael D.; Hallick, Richard B.

CS Department of Biochemistry and Molecular Biophysics, University of Arizona, Tucson, AZ, 85721, USA

SO Current Genetics (2001), 39(1), 49-60 CODEN: CUGED5; ISSN: 0172-8083

PB Springer-Verlag

DT Journal

LA English

AB The Escherichia coli aadA gene product, which confers resistance to spectinomycin and streptomycin, has been widely used as a dominant selectable marker for chloroplast transformation of Chlamydomonas and tobacco. An aadA expression in Euglena gracilis chloroplasts by replacing the Chlamydomonas promoter and 3' untranslated region (UTR) with the E. gracilis psbA promoter and 3' UTR. Transgenic DNA was introduced into E. gracilis chloroplasts by biolistic transformation. Streptomycin- and spectinomycin-resistant colonies were obtained, which screened pos. for the presence of the transforming vector by PCR amplification. Although integration of the transforming DNA into the chloroplast genome was not detected, transforming DNA was stably maintained in the chloroplast as an episomal element during continuous selection on antibiotics. The aadA cassette was also inserted into a transformation vector which contained the independently expressed psbK operon from either E. gracilis or a closely related species, E. stellata. The psbK operon contained at least two group III introns and a group III twintron, was highly expressed, and was only 1.5 kb in length. In transgenic E. gracilis chloroplasts, a truncated E. stellata psbK operon was transcribed, and the resultant pre-mRNA was accurately spliced. This system should allow the first direct anal. of group II and group III intron-splicing mechanisms. In addn., it could prove useful in the study of many other Euglena transcription and processing events. OSC.G 15 THERE ARE 15 CAPLUS RECORDS THAT CITE THIS RECORD (15 CITINGS) REICNT 63 THERE ARE 63 CITED REFERENCES AVAILABLE

RE ONT 63 THERE ARE 63 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2011 ACS on STN AN 1998:293059 CAPLUS < LOGINID::20110428>> DN 129:91014 OREF 129:18647a,18650a TI Root-directed expression of alien genes in transgenic potato: sarcotoxin and qus

AU Mahler-Slasky, Yael; Galili, Shmuel; Perl, Avihai; Aly, Radi; Wolf, Shmuel; Aviv, Dvora; Galun, Esra

CS Department of Plant Genetics, The Weizmann Institute of Science, Rehovot, Israel

SO Basic Life Sciences (1997), 65(Biology of Root Formation and Development), 187-192 CODEN: BLFSBY; ISSN: 0090-5542

PB Plenum Publishing Corp.

DT Journal

LA English

AB Bacterial pathogens of potato, e.g. Pseudomonas solanacearum, are known to infect potato roots causing severe losses in relatively warm climates. Our intention was to express, in potato roots, a bactericidal peptide that was identified in the larvae of the flesh fly (Sarcophaga peregrina) by Natori and assocs. in 1977. The cDNA coding for this peptide was subsequently isolated and it was termed sarcotoxin 1A (sarco) by these investigators. The resp. protein was also characterized and the mature proteins mass is about 5kDa. We used the Tob promoter that is root specific, to direct sarco expression in roots. In parallel we used the Gus gene as a reporter gene for this (Tob) promoter activity. Thus, two constructions of fusion genes were made. In one we inserted Tob up-stream of sarco and in the second Tob was inserted up-stream of Gus. In both cases the coding region was followed by a terminator. Both * * * transformation* * * ^{*} *** cassettes*** contained also a kanamycin (kana) resistance gene (nptII) that was inserted as

a selective marker under the 35S (CaMV) promoter. Agrobacterium-mediated genetic transformation was performed with potato tuber disks. Five potato cultivars and breeding lines were used: Desiree, Achirana INTA, LT-9, TS-10, TS-15. Potato plants that regenerated from Agrobacterium infected tuber-disks and rooted on selective medium were regarded putative transformants and were further analyzed. We found that putative transformants that resulted from transformation of a vector that contained Gus driven by Tob, indeed expressed the reporter gene in their roots. This verified the potency and specificity of the chimeric genes in the transformation vector. Polyclonal anti-sarco antibodies were produced and used to evaluate the expression of sarco in the putative transgenic potato plants. Preliminary western blot assays indicated that indeed the roots of some of the plants that were transformed with the chimeric-gene that contained the Tob promoter and the sarco cDNA, showed bands that reacted with the anti-sarco antibodies.

RE.ONT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2011 ACS on STN AN 1997:568296 CAPLUS < LOGINI D::20110428>> DN 127:230348

OREF 127:44819a,44822a

TI Recombinant expression cassettes for transformation of plant or other eukaryotes and regulation of gene expression in eukaryotes

IN Teasdale, Robert Dixon; Mouradov, Aidyn; Southerton, Simon George; Sawbridge, Timothy Ivor

PA Forbio Research Pty. Ltd., Australia; Teasdale, Robert Dixon; Mouradov, Aidyn; Southerton, Simon George; Sawbridge, Timothy Ivor

SO PCT Int. Appl., 87 pp. CODEN: PIXXD2

DT Patent

LA English
FAN: CNT 1 PATENT NO. KIND DATE
APPLICATION NO. DATE ------

PI WO 9730162 A1 19970821 WO 1997-AU89 19970219 W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, DK, EE, ES, FI, GB, GE, HU, IL, IS, CH, CN, CU, CZ, DE, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GR. GA, GN, ML, MR, NE, SN, TD, TG CA 2259456 19970821 CA 1997-2259456 19970219 AU 9717132 19970902 AU 1997-17132 19970219 EP 882133 A1 19981209 EP 1997-904302 19970219 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO CN 1216066 A 19990505 CN 19970219 JP 2000504577 1997-193833 20000418 JP 1997-528833 19970219 NO 9803775 19981015 NO 1998-3775 19980818 PRAI AU 1996-8161 WO 1997-AU89 A 19960219 19970219

AB There is provided a method of regulating a eukaryotically active gene, comprising transforming a cell with a

transformation ***cassette*** expressing a modulator gene product regulating the eukaryotically gene or its product and a further gene product regulating said modulator gene or its product, the promoters of two of said genes, modulator gene and further genes being selected from inducible promoters and developmental promoters for the same or complementary tissues. The lethal gene expressing barnase, a RNase of B. amyloliquefaciens, is placed under the control of a tissue specific promoter, such as those derived from PrMADS1, 2 or 3 of Pinus radiata or EGM1, 2 or 3 of Eucalyptus grandis. The same tissue specific promoter is used to express Laciq gene, a repressor for barnase (barstar) being promoted by a modified 35S RNA CaMV promoter including the lac operon. The cassette is used to transform plant cells for regeneration into plants expressing the barnase in the target tissues with improved specificity and reduced promoter leakage.

OSC.G. 8 THERE ARE 8 CAPLUS RECORDS THAT CITE
THIS RECORD (8 CITINGS)
RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE
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RE FORMAT

L7 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2011 ACS on STN AN 1995:346854 CAPLUS << LOGINID::20110428>> DN 122:98806

OREF 122:18495a,18498a

TI Transformation vectors that direct the integration of transforming DNA into the ribosomal DNA of a eukaryotic host IN Jacobs, Eric

PA Transgene S.A., Fr.

SO PCT Int. Appl., 35 pp. CODEN: PIXXD2

DT Patent

IA French

FAN. CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE -------

PI WO 9424300 A1 19941027 WO 1994-FR419 19940414 W: AU, CA, JP, US RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE FR 2703996 A1 19941021 FR 1993-4530 19930416 FR 2703996 B1 19950721 CA 2160697 A1 19941027 CA 1994-2160697 19940414 AU 9465719 19941108 AU 1994-65719 19940414 AU 686156 B2 19980205 EP 694072 A1 19960131 EP 1994-913647 19940414 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE JP 08508878 19960924 JP 1994-522836 19940414 US 6346414 B1 20020212 US 1995-532657 19951016 PRAL FR 1993-4530 A 19930416 WO 1994-FR419 W 19940414 ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

AB Transposition cassettes that preferably integrate into the ribosomal DNA of a eukaryotic host and based on a eukaryotic transposable element are described for use in gene therapy. The vectors carrying these cassettes also carry all the functions necessary for integration. The construction of a cassette for integration of transforming DNA into the human 28 S rRNA gene using the mobile intron 3 of the Carolina strain of Physarum polycephalum is demonstrated. This cassette was then introduced into an adenovirus that also carried an expression cassette for the P. polycephalum mobility endonuclease I-Ppo-I. A neomycin resistance marker was also included in the constructs.

OSC.G 4 THERE ARE 4 CAPLUS RECORDS THAT CITE

OSC.G 4 THERE ARE 4 CAPLUS RECORDS THAT CITE THIS RECORD (4 CITINGS)

RE.ONT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2011 ACS on STN AN 1994:319682 CAPLUS << LOGINID::20110428>> DN 120:319682

OREF 120:56129a,56132a

TI Transgenic plants with altered starch productivity
IN Keeling, Peter Lewis; Lomako, Joseph; Geowar-Singh,
Dave; Singletary, George William; Whelan, William Joseph
PA Zeneca Ltd., UK

SO PCT Int. Appl., 84 pp. CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE

A2 19940303 WO 1993-GB1821 PI WO 9404693 19930826 WO 9404693 A3 19940331 W: AU, BB, BG, BR, BY, CA, CZ, FI, HU, JP, KP, KR, KZ, LK, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SK, UA, US, VN RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG AU A 19940315 AU 1993-49707 19930826 AU 685065 B2 19980115 EP 658208 A1 19950621 EP 1994-908157 19930826 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE A 19990112 US 1995-392816 US 5859333 19951218

PRAI GB 1992-18185 A 19920826 WO 1993-GB1821 W 19930826

AB Plants with an altered starch synthesizing ability are produced by incorporating into the genome of the plant groreq.1 donor gene encoding a starch primer such as amylogenin, glycogenin, or protoglycogenin. The donor gene can be inserted into the host genome in the sense or antisense orientation. The recipient plant may be selected from Granimeae and Zea mays. The partially cloned amylogenin cDNA of B73 maize is disclosed and a procedure for the

construction of ***transformation*** ***cassette***
described.
OSC.G 10 THERE ARE 10 CAPLUS RECORDS THAT CITE
THIS RECORD (10 CITINGS)
RE ONT 1 THERE ARE 1 CITED REFERENCES AVAILABLE
FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE
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L3 12 S L2 NOT 2010/PY
L4 9 S L3 NOT 2009/PY
L5 6 S L4 NOT 2008/PY
L6 6 S L5 NOT 2007/PY
L7 6 S L6 NOT 2006/PY

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